Attro DAVID Schreiber

ACCESS DB# 13/12/11

## SEARCH REQUEST FORM

# Scientific and Technical Information Center

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Requester's Full Name:	NC ZARA	Examiner #: 7-75/2 Date: 10/5-/04	
Art Unit: /635 Phone N	lumber 20 2-0 76	Examiner #: $775/2$ Date: $10/5-0$ 4  Serial Number: $09/9/5$ , $3/4$	
Mail Box and Bidg/Room Location	:	uits Format Preferred (circle): HAPER DISK E-MA	ιiL
f more than one search is subm	itted, please prioritiz	ze searches in order of need.	**
include the elected species or structures, ke	eywords, synonyms, acror that may have a special m	as specifically as possible the subject matter to be searched.  nyms, and registry numbers, and combine with the concept or eaning. Give examples or relevant citations, authors, etc, if d abstract.	
Title of Invention: AS	modrof	1756	
Inventors (please provide full names):	n Bui	tler et ol.	
Earliest Priority Filing Date:	7126/01.	<u> </u>	
*For Sequence Searches Only* Please includ appropriate serial number.	le all pertinent information	(parent, child, divisional, or issued patent numbers) along with the	
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Searcher Phone #: 272 25 26	AA Sequence (#)		
learcher Location: Renstn Eal A6		Questel/Orbit	
Date Searcher Picked Up:	Bibliographic		
Date Completed: 1013	Litigation	Lexis/Nexis	
Searcher Prep & Review Time:	Fulltext	Sequence Systems can plag ~ c	
Clerical Prep Time:	Patent Family	WWW/Internet	
Online Time:	Other	Other (specify)	

PTO-1590 (8-01)

Attro DAvid Schreiber

Access	DB#	

### **SEARCH REQUEST FORM**

### Scientific and Technical Information Center

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Requester's Full Name: #			te: 10/8/04				
Art Unit: 1635 Phone N	Number 20 20 20	Serial Number: 09/9	15, 81 V				
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f more than one search is submitted, please prioritize searches in order of need.							
**************************************							
Include the elected species or structures, kutility of the invention. Define any terms known. Please attach a copy of the cover:	ceywords, synonyms, acro that may have a special n	onyms, and registry numbers, and combine eaning. Give examples or relevant cita	ne with the concept or				
Title of Invention: AS							
Inventors (please provide full names):	n Bu	theretal.					
Earliest Priority Filing Date:	7/26/01		•				
*For Sequence Searches Only* Please inclu appropriate serial number.	de all pertinent information	(parent, child, divisional, or issued patent	numbers) along with the				
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Date Completed:	Litigation	Lexis/Nexis					
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Clerical Prep Time:	Patent Family	WWW/Internet					
Online Time:	Other	_ Other (specify)_					

PTO-1590 (8-01)

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1 (bases 1 to 20)
Holly,R.D. and Foster,D.C.
Methods for producing thrombin
Patent: US 5476777-A 5 19-DEC-1995;
Location/Qualifiers
                                                                                                                                                                                                                                                                              16;
                                                                                                                                                                                                                                                                                                                                                     Unclassified.

1 (bases 1 to 20)
Holly,R.D. and Foster,D.C.
Methods for producing thrombin
Patent: US 5502034-A 5 26-MAR-1996;
Location/Qualifiers
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Sequence 5 from patent US 5502034.
I19059
I19059.1 GI:1599414
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I16620
I16620.1 GI:1251528
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                               Holly R.D. and Foster, D.C.
Holly R.D. and Foster, D.C.
Methods for producing thrombin
Patent: US 5527692-A 5 18-UUN-1996;
Location/Qualifiers
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larity 100.0%; Pred. No.
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/mol_type="unassigned DNA"
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/mol_type="unassigned DNA"
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Novel human protein kinases and protein kinase-like enzymes Patent: WO 0138503-A 218 31-WAY-2001; Sugen, Inc. (US)
                                                                                                                                                                                                                                                                                                                                                                                                Sequence 218 from Patent
AX166727
AX166727.1 GI:14547002
                                                                                                                                                                                                                                                                                                                                                            Homo sapiens (human)
                      Sequence 1478 from Patent AX531969
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Mammalia; Eutheria; Primates;
            AX531969.1 GI:25255707
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nilarity 89.5%;
Conservative
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/mol_type="unassigned DN
/db_xref="taxon:9606"
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Pred. No. 36;
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Catarrhini; Hominidae;
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REFERENCE AUTHORS TITLE JOURNAL

Shannon, M.

FEATURES

Human posh-like protein 1 Patent: EP 123051-A 1478 11-SEP-2002; Aeomica, Inc. (US) Location/Qualifiers

/organism="Homo sapiens" /mol\_type="unassigned DNA" /db\_xref="taxon:9606"

ORGANISM

Homo sapiens (human)

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

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CAGCAGCCTGATAAAGAGCCACGG

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Query Match
Best Local
        Similarity
       0.7%;
87.5%;
0;
        Score 19.2;
Pred. No. 1
Mismatches
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             Length 24;
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14-MAY-2003

(first entry)

RESULT 138 ABZ82615

ABZ82615

standard; DNA; 19

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ABZ82615,

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19

Matches

Best Local Similarity
Matches 19; Conserv

Conservative

0;

100.0%;

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Score 19; Pred. No. Mismatches

DB 1;

Length 19; Indels

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2114 GGACAGGACAGTGAGGAGC 2132

Query Match

Sequence 24 BP; 8 A;

7 C; 7

G; 2 T; 0 U; 0 Other;

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2134 CAGCAGCCTGATAAAGTCCAACGG

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RESULT 136
AB189232
XX AB1892
XX AB1892
XX AB1892
XX AB1892
XX AB1892
XX AB1892
XX Captux
XX Human;
XX Human;
XX Infect
XX Synthm
XX Synthm
XX WO200]
XX WO200]
XX How wo2
                                                                 The present invention describes a method (MI) for designing capture CC oligonucleotide probes (II) will hybridise with little mismatch, where CC (I) have melting temperatures within a narrow range. The method is useful for detecting infectious diseases caused by bacterial infectious agents CC e.g. Salmonella, Listeria monocytogenes and Haemophilus influenza, fungal infectious agents e.g. Cryptococcus neoformans, Candida albicans and CC Aspergillus fumigautus, viruses e.g. T-cell lymphocytotrophis cirus, CC Epstein-Barr virus and polio virus, and parasitic infectious agents CC selected from Onchoverva volvulus, Entamoeba histolytica and Dracunculus CC medinesis. The method is also useful for detecting genetic diseases such as 21 hydroxylase deficiency, Turner Syndrome and obesity defects. CC Detecting cancer involving oncogenes, tumour suppressor genes, or genes involved in DNA amplification, replication, recombination or repair, the cancer is specifically associated with a gene selected from BRCA1 gene, 53 gene, human papillomavirus types 16 and 18 and liver cancers. The centhod is also used for environmental monitoring, forensics and the food and feed industry, detecting comprises scanning (using e.g. a scanning clectron microscope and infrared microscope) the support at the comparison of the oligonucleotide probe sets occurred and correlating (using a computer) identified ligation to a present oligonucleotide sequences. ABI97546 represent oligonucleotide sequences. ABI9774 to of the present invention
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Designing capture oligonucleotide probes complementary oligonucleotides hybridize
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RESULT 137
AAT43117/c
ID AAT431
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XX DT 05-SEP
XX Immort
KW Immort
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KW medium
KW amplif
XX AT610
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                                                                  The invention relates to new immortalised cell lines derived from pre-
adipocytes containing an immortalising fragment of a viral oncogene. The
immortalised adipocytes are used to identify substances able to regulate
lipolysis and/or thermogenesis (potential therapeutic agents for treating
diabetes and obseity). The cell lines have the advantage that they can be
maintained in long term oulture (contrast primary cultures of adipocytes)
without loss of characteristic markers or ability to differentiate. The
immortalised pre-adipocytes differentiate into mature adipocytes when
placed in a medium containing insulin and dexamethasone. The primers
AAT43098-19 are used to amplify marker genes to verify differentiation of
the pre-adipocytes into mature adipocytes. Purimers AAT43116-7 were used
to amplify a 286 by region of the gene encoding a hormone sensitive
lipase, a marker for mature "brown" adipocytes
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       Sequence 19
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9 C; 3 G; 6 T; 0 U; 0 Other;
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AR139573
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Strosberg, A. Donny. and Zilberfarb, V.
Immortalized cell lines from human adipose tissue, process
preparing same and applications thereof
Patent: US 6071747-A 20 06-JUN-2000;
   1 (bases 1 to 21 Keating, M.T. and Mutations in and
                                                 Unknown
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Sequence 20 from
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IMMORTALISED CELL LINES FROM HUMAN ADIPOSE
PREPARING SAME AND APPLICATIONS THEREOF
Patent: WO 9634100-A 20 31-OCT-1996;
CENTRE NAT RECH SCIENT (FR)
Other publication FR 2733513 961031.
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/mol_type="unassigned DNA"
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/mol_type="unassigned DNA"
/db_xref="taxon:32644"
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UNIVERSITY OF UTAH RESEARCH FOUNDATION
OS HOMO sapiens (human)
PN JP 2002521065-A/88
PD 16-JUL-2002
PP 20-JUL-1999 UP 2000562554
PF 20-JUL-1999 UP 2000562554
PR 27-JUL-1998 US 09/122847,06-JAN-1999 US 09/226012 PI
MARK T KEATING, IGOR SPLAWSKI
C12N15/09,A01K67/027,C07K14/47,C07K16/18,C12N1/15,C12N1/19, PC
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Mutations
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 21)
Keating,M.T. and Splawski,I.
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                                                         Conservative
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                                                                                                                                                                                                                                          syndrome gene
                                                                                                                /organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
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PAT 03-MAR-1998